By order of the Board of Governors of the Federal Reserve System, January 18, 1995.

William W. Wiles,

Secretary of the Board.

[FR Doc. 95–1786 Filed 1–25–95; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Chapter I

[Docket No. 95D–0014]

Draft Proposed Regulations on Quality Standards and Certification Requirements for Mammography Facilities; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability of draft proposed regulations.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of draft proposed regulations prepared by the Center for Devices and Radiological Health (CDRH) to implement the Mammography Quality Standards Act of 1992 (MQSA). The drafts contain minimum quality standards for mammography facilities in the following areas: Personnel standards, medical outcomes audits, medical records and mammography reports, quality assurance, mobile mammography, accreditation bodies, implant imaging, mammography equipment, variances, consumer complaint mechanism, and quality control.

DATES: Written comments by April 11, 1995.

ADDRESSES: Submit written requests for single copies of the draft regulations to the Center for Devices and Radiological Health (HFZ–240), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850. Send two self-addressed adhesive labels to assist that office in processing your requests. Submit written comments on the draft proposed regulations to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857. Requests and comments should be identified with the docket number found in brackets in the heading of this document. A copy of the draft proposed regulations and received comments are available for public examination in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Charles K. Showalter, Center for Devices and Radiological Health, Center for Devices and Radiological Health (HFZ–240), 9200 Corporate Blvd., Rockville, MD 20850, 301–594–3311.

SUPPLEMENTARY INFORMATION: FDA is announcing the availability of draft proposed regulations regarding quality standards and certification requirements for mammography facilities in the following areas: Personnel standards, medical outcomes audits, medical records and mammography reports, quality assurance, mobile mammography, accreditation bodies, implant imaging, mammography equipment, variances, consumer complaint mechanism, and quality control. These proposed regulations are being developed to implement the MQSA (Pub. L. 102–539), which was enacted to establish quality standards for mammography. The MQSA requires that, to provide mammography services legally after October 1, 1994, all facilities, except facilities of the Department of Veterans Affairs, must be both accredited by an approved accrediting body and certified by the Secretary of Health and Human Services (the Secretary). The authority to approve accrediting bodies and to certify facilities has been delegated by the Secretary to FDA.

Pursuant to authorization from Congress, FDA promulgated interim regulations to ensure that mammography facilities meet minimum quality standards. These regulations, which were published in the **Federal Register** on December 21, 1993 (58 FR 67558 and 58 FR 67565), and amended on September 30, 1994 (59 FR 49808), became effective on October 1, 1994, and will remain in effect until final regulations are promulgated.

FDA is currently developing proposed regulations for quality standards in various subject areas, including the 11 areas referenced above. Under the MQSA, FDA established a National Mammography Quality Assurance Advisory Committee (NMQAAC) to advise the agency on the appropriate level of quality standards for mammography facilities and accrediting bodies. Advanced drafts of proposed regulations are provided routinely to all members of the advisory committee for their advice and recommendation, and periodic public meetings of the advisory committee are being held.

An advisory committee meeting was held on January 23 through January 25, 1995. The meeting was announced in the **Federal Register** of December 21, 1994 (59 FR 65776). It was held at the Dupont Plaza Hotel, Embassy Room, 1500 New Hampshire Ave. NW., Washington, D.C.

In order to gather additional information on these particular topics, FDA decided to share the drafts of these proposed regulations with certain individuals who were invited as guests to the January 23 through 25, 1995, advisory committee meeting to enable them to provide comments at the meeting on the feasibility of efficient implementation of these draft proposed standards by the radiology community. These invited guests have particular expertise in one or more of the areas addressed by the draft proposals. The agency is publishing this notice in order to make the same draft documents available to the general public.

Although all members of the general public will have an opportunity to comment on the proposed regulations when they are published in the spring of 1995, interested persons who wish to comment on the draft proposals may submit written comments to the Dockets Management Branch (address above) on or before April 11, 1995. Two copies of any comments should be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. The draft proposed regulations and comments received may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

FDA will consider any comments received in developing final regulations.

Dated: January 23, 1995.

William K. Hubbard,

Interim Deputy Commissioner for Policy.

[FR Doc. 95–2006 Filed 1–23–95; 3:40 pm]

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DEPARTMENT OF JUSTICE

Office of Justice Programs

28 CFR Part 93

[OJP No. 1014]

RIN 1121–AA26

Drug Courts

AGENCY: Department of Justice, Office of Justice Programs.

ACTION: Proposed rule.

SUMMARY: This notice announces a proposed rule and requests comments on the Drug Court Program as authorized by Title V of the Violent Crime Control and Law Enforcement Act of 1994. This rule gives general